National specialized resources for RD in each country are collected by dedicated Orphanet National Teams. Information is collected from selected data sources and manually curated. The table below indicates in green the type of resources that will be collected in 2024 by each national team, according to team's priorisation and ressources. To know more about the sources that are surveyed in each country you can consult the National websites available here to know more about the procedures for data collection click here https://www.orpha.net/en/other-information/procedures

If you want to get in touch with the Orphanet National Team contact forms are available here: https://www.orpha.net/consor/cgi-bin/Directory_Contact.php?lng=EN

Country	Expert Centres	Patient Organisations	Laboratories/Diagnostic Tests	Research Projects	Clinical Trials	Registries	Biobanks	infrastructures/Platforms
AT	🟩	🟩	🟥 *	🟩	🟩	🟥	🟥	🟥
BE	🟩	🟩	🟩	🟩	🟩	🟩	🟩	🟩
BG	🟩	🟩	🟥	🟥	🟥	🟥	🟥	🟥
CZ	🟩	🟩	🟥	🟥	🟥	🟥	🟥	🟥
CH	🟩	🟩	🟥 *	🟩	🟩	🟩	🟩	🟩
DE	🟩	🟩	🟥 *	🟩	🟩	🟩	🟩	🟩
EE	🟩	🟩	🟥	🟩	🟩	🟩	🟩	
ES	🟩	🟩	🟩 EQAS programme	🟩	🟩	🟩	🟩	🟩
FI	🟩	🟩	🟥	🟩	🟩	🟩	🟩	🟩
FR	🟩	🟩	🟥	🟩	🟩	🟩	🟩	🟩
GE	🟨	🟨	🟨	🟨	🟨	🟨	🟨	🟨
GR	🟩	🟩	🟥	🟥	🟥	🟥	🟥	🟥
HR	🟨	🟨	🟨	🟨	🟨	🟨	🟨	🟨
IE	🟩	🟩	🟥 *	🟩	🟩	🟩	🟩	🟥 *
IL	🟨	🟨	🟨	🟨	🟨	🟨	🟨	🟨
IT	🟩	🟩	🟥	🟩	🟩	🟩	🟩	🟩
JP	🟨	🟨	🟨	🟨	🟨	🟨	🟨	🟨
KZ	🟨	🟨	🟨	🟨	🟨	🟨	🟨	🟨
LT	🟩	🟩	🟩	🟥	🟥	🟥	🟥	🟥
LU	🟩	🟩	🟥	🟥	🟥	🟥	🟥	🟥
LV	🟩	🟩	🟥	🟥	🟥	🟥	🟥	🟥
MK	🟩	🟩	🟩	🟥	🟥	🟥	🟥	🟥
NL	🟩	🟩	🟩	🟥 *	🟩	🟥 *	🟥 *	🟥 *
NO	🟩	🟩	🟥 only labs (for tests: https://www.genetikkportalen.no/)	🟩	🟩	🟩	🟩	🟩
PL	🟨	🟨	🟨	🟨	🟨	🟨	🟨	🟨
PT	🟩	🟩	🟥	🟥	🟥	🟩	🟥	🟥
RO	🟩	🟩	🟥 *	🟩	🟥 *	🟥 *	🟥 *	🟥 *

RS	🟩	🟩	🟥	🟩	🟥	🟥	🟥	🟥
SE	🟩	🟩	🟥 *	🟩	🟩	🟩	🟥	🟥
SI	🟩	🟩	🟩	🟩	🟥	🟥	🟥	🟥
UK	**Observer**							
RU	**Temporary Suspension**							
TR	🟨	🟨	🟨	🟨	🟨	🟨	🟨	🟨

🟩	Info will be collected in 2024
🟥	Info will not be collected in 2024
🟥 *	Info will only be collected if requested by professionals
🟨	no info available regarding 2024 priorisation